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54 **Process for preparing polymers by anionic polymerization.**

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Publications Ltd, London, GB;**

73 Proprietor: **THE DOW CHEMICAL COMPANY**
2030 Dow Center
Abbott Road
P.O. Box 1967
Midland Michigan 48640-1967(US)

72 Inventor: **Marchand, Gary R.**
4028 Lassen Drive
Baton Rouge, Louisiana 70814(US)
Inventor: **Walther, Brian W.**
9378 Tasmania Avenue
Baton Rouge, Louisiana 70810(US)
Inventor: **Rose, Warren Ronald**
1751 Twisted Oak Lane
Baton Rouge, Louisiana 70810(US)

74 Representative: **Sternagel, Hans-Günther, Dr.**
et al
Patentanwälte Dr. Michael Hann
Dr. H.-G. Sternagel
Sander Aue 30
D-51465 Bergisch Gladbach (DE)

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Description

The present invention relates to a process for preparing polymers by means of anionic polymerization. More particularly the present invention relates to a new method for the termination of such anionic polymerization reactions or for treating coupled reaction products of such polymerization reactions.

It is previously known in the art to prepare polymers of anionically polymerizable monomers, especially monovinylidene aromatic monomers and alkadiene monomers via anionic polymerization. Highly useful polymers are prepared in the form of block copolymers containing one or more blocks of a monovinylidene aromatic polymer and one or more blocks of an alkadiene polymer. Suitably, such polymers are prepared in an organic solvent and are usefully employed in adhesive formulations, as modifiers for thermoplastic resins and asphalt or bituminous compositions or in solvent containing cements or mastic formulations. The polymers containing residual unsaturation may be hydrogenated if desired to produce products having improved weathering resistance.

The initiators used in such anionic polymerizations are typically very strong bases. Examples are the alkali metal organyl compounds, particularly alkali metal alkyls, especially sodium and lithium alkyls. After polymerization is complete, the terminal monomer group of the living polymer anion must be terminated. Termination may occur through a coupling reaction by means of a coupling agent or, in the event a coupled compound is not desired, by the use of suitable proton donating agents such as an organic alcohol, ammonia, amines or even water to terminate the living anion. However, the remnant formed by this termination is itself a basic species. In the industrial preparation of polymers via anionic polymerization, especially the preparation of block copolymers of monovinylidene aromatic monomers and alkadienes, it is often desirable to include an antioxidant in the polymeric syrup to prevent oxidative and mechanical degradation of the polymer during devolatilization and finishing. However, many of the suitable antioxidants employed in such formulations are sensitive under basic conditions. That is, such antioxidants react with basic species thereby forming undesirable products. Alternatively under basic conditions the antioxidant may be inhibited in its ability to prevent oxidative degradation of the resulting polymer.

Consequently, products which have been neutralized by the use of organic alcohols, especially such products further containing an antioxidant, have been found to be lacking in both color retention and in aging stability. Such polymeric products have been found to change in melt viscosity after further mechanical and thermal treatment and to be marked by an absence of clarity as a result of increased haze. In addition such polymeric products tend to have increased yellowing. Also, physical properties, such as ultimate tensile strength, are adversely affected.

In U.S. Patent 4,415,695 it is proposed to employ boric acid as a terminating agent in an anionic polymerisation. Disadvantageously when boric acid is employed as a terminating agent in the preparation of a block copolymer of a monovinylidene aromatic monomer and an alkadiene, the resulting product still possesses an undesirable change in melt viscosity upon thermal aging.

US-A-2,543,440 discloses a process in which the reaction product of the anionic polymerization of conjugated dienes is contacted with a solution or mixture of water and a low molecular weight alcohol to which has been added a minor portion of an acid, preferably a mineral acid. In one embodiment of this process the reaction product is first contacted with isopropanol and thereafter with an aqueous solution of sulfuric acid and sodium sulfate.

GB-A-776,682 teaches treating the reaction product of the anionic polymerization of butadiene or copolymerization of butadiene and styrene with aqueous sulfuric acid containing more than 80 and less than 96% H_2SO_4 in an amount sufficient to convert the alkali metal compound quantitatively to its bisulphate.

In accordance with WPIL No. 88-126,730, Derwent Publications, orthophosphoric acid having a concentration of from 40 to 84 wt.% is added to the aforesaid reaction products, followed by centrifuging and/or extracting the metal salts formed with water. All the said prior art processes, though terminating the polymerization reaction effectively, do not yield a polymerizate with properties which are satisfactory in all respects.

It would be desirable if there were provided an improved process for preparing polymers by means of anionic polymerization techniques employing a terminating agent which does not adversely affect the polymer properties, such as ultimate tensile strength.

It would also be desirable if there were provided an improved technique for treating terminated reaction products prepared by anionic polymerization that allows for the preparation of polymers having improved clarity and other polymer properties.

It would be desirable if there were provided an improved technique for treating coupled reaction products prepared by anionic polymerization that allows for the preparation of polymers having improved

clarity and other polymer properties.

Finally, it would be desirable if there were provided adhesives containing as one component the improved block copolymers of monovinylidene aromatic monomers and alkadienes prepared according to the present invention.

5 According to the present invention there is now provided a process for preparing polymers by means of the anionic polymerization of polymerizable monomers the steps of the process comprising:

- (a) contacting one or more anionically polymerizable monomers selected from monovinylidene aromatic monomers and alkadienes with a lithium containing initiator under anionic polymerization conditions;
- (b) terminating the polymerization by contacting the reaction mixture resulting from step (a) with a proton
10 donating agent resulting in a lithium containing basic species;
- (c) neutralizing the lithium containing basic species by contacting the reaction mixture from step (b) with from 0.01 to 10 equivalents of phosphoric acid, based on the lithium containing initiator; and
- (d) recovering the resulting polymer.

Also included within the scope of the present invention is a polymeric product prepared by anionic
15 polymerization according to the foregoing techniques as well as formulated products such as adhesives prepared therefrom. In a particular embodiment such a polymerization product also contains residual quantities of a lithium compound which is the remnant of the polymerization initiator and an amount of phosphoric acid sufficient to neutralize the lithium compound.

Processes for the anionic polymerization of monomers are well known in the art. Initial work utilizing
20 sodium initiators such as sodium naphthalene (J. Amer. Chem. Soc., 78, 2656, 1956) was later followed by lithium containing initiators such as secondary butyllithium (U.S. Patents 3,321,635 and 3,265,765). More recent process improvements have led to precise control of the polymerization to produce polymers having a wide variety of physical properties. Examples include EP 210,677 which discloses an adiabatic polymerization of styrene in cyclohexane followed by addition of butadiene monomer. Additional anionic
25 techniques include hybrid Ziegler-Natta/anionic methods disclosed in USP 4,480,075; the use of Lewis bases to control diene vinyl content (USP 4,530,985); and various hydrogenation techniques to provide saturated polymeric products (USP 4,595,749, 4,035,445).

In addition to monofunctional initiators such as the aforementioned lithium alkyls, there have been proposed certain difunctional lithium-containing initiators suitable for use in the direct preparation of block
30 copolymers of dienes and monovinylidene aromatic monomers. Such difunctional initiators are disclosed in U.S. Patents 3,660,536; 3,776,893; 3,954,894; 4,172,190; 4,196,153; 4,200,718; 4,205,016; 4,431,777; 4,427,837; and 4,614,768. Preferred difunctional initiators are, for example, 1,3-phenylene-bis(3-methyl-1-phenyl pentylidene)bis-lithium, the isomeric methylphenyl substituted derivatives such as 1,3-phenylene-bis-(3-methyl-1-(2-methylphenyl)-pentylidene, 1,3-phenylene bis(3-methyl-1-(4-methylphenyl)pentylidene, and
35 mixtures thereof.

Suitable monomers are those in the class of anionically polymerizable monomers such as alkadienes, monovinylidenes, alkyl acrylates, acrylamides, acrylonitrile, vinylsilanes, arylsilanes, vinylketones, vinylpyridines, isocyanates, diisocyanates, oxides such as ethylene oxide, lactams and siloxanes.

Preferred polymerizable monomers for use according to the present invention include the well known
40 alkadienes especially butadiene and isoprene, and monovinylidene aromatic monomers, especially styrene and α -methylstyrene as well as ring alkyl substituted derivatives thereof. A preferred monovinylidene aromatic monomer is styrene.

The amount of phosphoric acid added according to the present invention should be sufficient to neutralize the polymer anion contained in the reaction mixture. Suitably the equivalent ratio of neutralizing
45 agent added to the reaction mixture based on initial initiator added (i.e. the ratio of equivalents agent/equivalents initiator) is from 0.01 to 10.0, preferably from 0.75 to 2.00. After addition of the neutralizing agent, antioxidant package and additional optional ingredients, the solvent is removed by devolatilization or other suitable technique and the resulting polymer recovered.

In accordance with the present invention the remnant initiator is first terminated according to conven-
50 tional techniques utilizing a proton donating terminating agent and the resulting basic species then neutralized. While it may be possible to obtain thorough incorporation of the phosphoric acid if the process is accomplished at the time of termination, acceptable results are obtained if the polymer product is blended with the neutralizing agent at a later time. Suitably the blending step may be accomplished by use of a ribbon blender, extruder, or other suitable malaxing device. Amounts of neutralizing additive incor-
55 porated into the polymer generally are less than 1.0 percent based on total polymer weight.

As previously mentioned, antioxidants may be usefully incorporated in the present products. Suitable antioxidants include those compositions previously known and utilized in the art for antioxidant purposes. Examples include hindered phenolic, phosphite, phosphonite, or phosphate type antioxidants. Other ad-

ditives such as, for example, extrusion aids, UV light stabilizers and viscosity modifiers may be incorporated as desired.

Formulated products are readily prepared from the products of the present invention as is well known in the art. For example, block copolymers of monovinylidene aromatic monomers and alkadienes are usefully combined with tackifiers such as hydrogenated terpenes, low molecular weight polypiperilidenes, and other suitable compounds to form an adhesive composition.

Having described the invention the following examples are provided as further illustrative of the invention and are not to be construed as limiting.

10 Examples 1 to 6

A five gallon (19 liters) lab reactor was charged with 12.1 kg of cyclohexane, 1.70 kg of isoprene and the solution was heated to a temperature of 45 °C. Then 19 g of a 0.0282 M/l cyclohexane solution of 1,3-phenylene-bis(3-methyl-1-phenylpentylidene)bis-lithium was added to the reaction solution to remove impurities. Next 380.6 g of the same 0.0282 M/l cyclohexane solution of the bis-lithium compound was added to initiate polymerization. After all the isoprene was polymerized and the reaction mixture had cooled to 60 degrees, 276.4 g of styrene monomer were added to the reaction solution. After all the styrene monomer was polymerized, 1.84 g of neat isopropanol was added to the reaction solution. The resulting polymer syrup, containing a styrene-isoprene-styrene triblock copolymer, was then separated into portions for use in the neutralization and stabilization experiments. Six approximately 1 liter portions of the above polymer syrup were weighed. The resulting polymer syrup was calculated to contain 2.01×10^{-6} moles Li/g solution.

The six portions were treated as follows. The amount of neutralizing agent used, if any, was calculated by assuming one mole basic species/mole Li. Phosphoric acid was treated as a one proton donor. Sulfuric acid was treated as a two proton donor. The acid concentrations were obtained from calculations based on reported acid purities, densities, and molecular weights. Then 0.3 weight percent of 2,2-bis[[3-[3,5-bis(1,1-dimethyl-ethyl)-4-hydroxyphenyl]-1-oxopropoxy]methyl]-1,3-propanediyl 3,5-bis(1,1-dimethylethyl)-4-hydroxybenzene propanoate (Irganox® 1010), and 0.3 weight percent of trisnonylphenyl phosphate stabilizer were added.

30 Devolatilization

The neutralized and stabilized polymer syrups were devolatilized at 90 °C and at 8-25 psi (55-172 kPa) vacuum for 2.5 hr. The polymers are cooled under about 30 psi (207 kPa) vacuum for at least 1 hr.

35 Melt Viscosity Determinations

Melt viscosity determinations were made using a conventional melt indexer. The melt index determinations were made at 200 °C with a 5 kg weight and a 2.1 mm x 8 mm die (ASTM-D1238, condition 200/5.0). The results are shown in Table 1.

40 Ultimate Tensile Measurements

Strips of polymer cut from the devolatilized slab were compression molded and tensile specimens were die cut. The compression molding conditions were for 7.0 g sample in a 3" x 4.5" x 0.035" (76 mm x 114 mm x 0.9 mm) chase. The press temperature was 200 °C. Samples were pre-heated for 3.0 minutes under approximately 2000 kg/cm² pressure. The samples were then cured for 3 minutes at 18000 kg/cm² pressure and then cooled to ambient temperature from 200 °C at a rate of 30-50 °C/min. The sample plaques were then die cut using a NAEF® stamping press with an ASTM D 1822, type L die. Ultimate tensile strengths were measured on a Monsanto T-10 tensometer. Special elastomeric grips were required to prevent cutting of the sample by the clamping surfaces. The thickness of the sample was recorded to 0.0001" (0.0025 mm) using a micrometer and is inserted into the tensometer so as to attain a gauge length of 1.0" (2.54 cm). The samples were pulled at a cross head speed of 10 in/min (25.4 cm/min) and the tensile value was obtained at break. A summary of the data is contained in Table 1.

Roll Milling

Portions of the devolatilized polymer slabs were roll milled twice for 3.5 minutes at 155°C at a gap width of 0.028" (0.71 mm).

Tests on Roll Milled Samples

Melt index and in some cases ultimate tensile determinations were performed on the roll milled polymers. Melt index and tensile determinations followed the same procedure as for the previously disclosed slab materials.

Color and Haze Tests

A portion of the plaque made from roll milled polymer was tested for yellowness index and percent haze according to ASTM D 1925-70 and ASTM D 1003-61, respectively. A Hunterlab Inc. Tristimulus® colorimeter was used for both measurements. The instrument was standardized for diffuse transmittance. The data are located in Table I.

TABLE I

<u>Example</u>	<u>Neutralizer</u>	<u>Percent Change After Roll Milling</u>		<u>Yellow Index c</u>	<u>Trans. Haze c</u>
		<u>Melt Index a</u>	<u>Ultimate Tensile b</u>		
1	phosphoric acid	24%	6%	3.1	23%
Compa- rative:					
2	none	35%	32%	3.2	28%
3	stearic acid	43%	-	2.3	22%
4	sulfuric acid	14%	15%	2.0	22%
5	nitric acid	61%	48%	3.5	52%
6	hydrochloric acid	-	40%	5.0	74%

a = ((Final MFR/Initial MFR)-1)*100

b = (1-(Final Tensile/Initial Tensile))*100

c = Roll Milled Polymer

As may be seen by reference to Table I, the use of phosphoric acid to neutralize remnants of the lithium initiator resulted in a product having improved ultimate tensile strength after exposure to further processing conditions.

Example 7

A five gallon (19 liters) reactor was charged with 12.5 kg of cyclohexane, 285 g of styrene, and the solution was heated to a temperature of 45 °C. Then 70 g of a 0.295 M/l cyclohexane solution of sec-butyl lithium was added to initiate polymerization. After all the styrene was polymerized and the reaction mixture had cooled to 45 °C, 1565 g of butadiene monomer was added to the reaction solution. After all the butadiene monomer was polymerized and the reactor was cooled to 45 °C, 285 g of styrene was added. After all the styrene was polymerized, about 2 ml of isopropanol was added to quench the reaction. The resulting polymer syrup, containing a styrene-butadiene-styrene triblock copolymer, was then separated into portions for use in the neutralization and stabilization experiments. Two portions of approximately 2 liters of the above polymer syrup were weighed. The polymer syrup was calculated to contain 1.82×10^{-6} moles Li/g of solution.

The two portions were treated as follows. To one sample of polymer syrup no phosphoric acid was added. Phosphoric acid was added to the other sample in a ratio of one mole of acid to one mole of lithium. The acid concentration was obtained from a calculation based upon the reported acid purity, density, and molecular weight. 0.5 Weight percent of Irganox® 565 and 0.5 weight percent of Irgafos® 168 were then added. Devolatilization, melt viscosity determination, ultimate tensile measurement, and color and haze tests were performed as in Examples 1 and 2. The total time of roll milling was 20 minutes for this study. The data from these tests are located in Table II.

TABLE II

Example	Acid Ratio to Li	Percent Change After Roll Milling		Yellow Index c	Trans. Haze c
		Melt Index a	Ultimate Tensile b		
Comparative 7	0	64	41	3.1	13.3
	1.0	43	15	3.9	13.4

a = ((Final MFR/Initial MFR)-1)*100

b = (1-(Final Tensile/Initial Tensile))*100

c = Roll Milled Polymer

Claims

1. A process for preparing polymers by means of the anionic polymerization of polymerizable monomers, the steps of the process comprising:
 - (a) contacting one or more anionically polymerizable monomers selected from monovinylidene aromatic monomers and alkadienes with a lithium containing initiator under anionic polymerization conditions;
 - (b) terminating the polymerization by contacting the reaction mixture resulting from step (a) with a proton donating agent resulting in a lithium containing basic species;
 - (c) neutralizing the lithium containing basic species by contacting the reaction mixture from step (b) with from 0.01 to 10 equivalents of phosphoric acid, based on the lithium containing initiator; and
 - (d) recovering the resulting polymer.
2. The process as claimed in claim 1 wherein the proton donating terminating agent is an alcohol, ammonia, an amine or water.
3. The process as claimed in claim 1 or 2 wherein the amount of phosphoric acid added to the reaction mixture of step (a) is from 0.75 to 2.00 equivalents, based on the lithium containing initiator.
4. The process as claimed in any of claims 1 to 3 wherein the monovinylidene aromatic monomer is styrene and the alkadiene is isoprene or butadiene..
5. An adhesive formulation comprising a polymer prepared according to the process of any of claims 1 to 4.

Patentansprüche

1. Verfahren zur Herstellung von Polymeren mittels anionischer Polymerisation von polymerisierbaren Monomeren, bei dem man
 - (a) eines oder mehrere anionisch polymerisierbare Monomere, ausgewählt aus der Gruppe, bestehend aus monovinylidenaromatischen Monomeren und Alkadienen, mit einem Lithium enthaltenden Initiator unter den Bedingungen der anionischen Polymerisation in Berührung bringt;
 - (b) die Polymerisation beendet, indem man das Reaktionsgemisch der Stufe (a) mit einem Protonendonator in Berührung bringt, wodurch eine Lithium enthaltende basische Verbindung entsteht;
 - (c) die Lithium enthaltende basische Verbindung neutralisiert, indem man das Reaktionsgemisch der Stufe (b) mit 0,01 bis 10 Äquivalenten Phosphorsäure, bezogen auf den Lithium enthaltenden Initiator, in Berührung bringt; und
 - (d) das entstandene Polymere isoliert.
2. Verfahren nach Anspruch 1, wobei der die Reaktion beendende Protonendonator ein Alkohol, Ammoniak, ein Amin oder Wasser ist.
3. Verfahren nach den Ansprüchen 1 oder 2, wobei die Menge der dem Reaktionsgemisch der Stufe (b) zugesetzten Phosphorsäure von 0,75 bis 2,00 Äquivalente, bezogen auf den Lithium enthaltenden Initiator, beträgt.
4. Verfahren nach jedem der Ansprüche 1 bis 3, wobei das monovinylidenaromatische Monomere Styrol und das Alkadien Butadien oder Isopren ist.
5. Klebe- oder Haftmittel, enthaltend ein Polymeres, hergestellt nach einem der Ansprüche 1 bis 4.

Revendications

1. Procédé pour préparer des polymères au moyen d'une polymérisation anionique de monomères polymérisables, les étapes du procédé comprenant les opérations consistant:
 - (a) à mettre en contact un ou plusieurs monomères anioniquement polymérisables choisis parmi les monomères aromatiques monovinylidène et les alcadiènes avec un initiateur contenant du lithium dans des conditions de polymérisation anionique;
 - (b) à terminer la polymérisation en mettant en contact le mélange réactionnel résultant de l'étape (a) avec un agent donneur de proton aboutissant à des espèces basiques contenant du lithium;
 - (c) à neutraliser les espèces basiques contenant du lithium en mettant en contact le mélange réactionnel de l'étape (b) avec de 0,01 à 10 équivalents d'acide phosphorique, par rapport à l'initiateur contenant du lithium; et
 - (d) à récupérer le polymère résultant.
2. Procédé selon la revendication 1, dans lequel l'agent de terminaison donneur de proton est un alcool, de l'ammoniac, une amine ou de l'eau.
3. Procédé selon l'une des revendications 1 ou 2, dans lequel la quantité d'acide phosphorique ajoutée au mélange réactionnel de l'étape (a) est de 0,75 à 2,00 équivalents, par rapport à l'initiateur contenant du lithium.
4. Procédé selon l'une quelconque des revendications 1 à 3, dans lequel le monomère aromatique monovinylidène est le styrène, et l'alcadiène est l'isoprène ou le butadiène.
5. Formulation d'adhésif comprenant un polymère préparé selon le procédé de l'une quelconque des revendications 1 à 4.